

IAP5 Rec'd PCT/PTO 27 SEP 2006
DESCRIPTION

HEAT-RESISTANT PHOTSENSITIVE RESIN COMPOSITION, METHOD FOR
FORMING PATTERN USING THE COMPOSITION, AND ELECTRONIC PART

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TECHNICAL FIELD

The present invention relates to a heat-resistant
photosensitive resin composition, and, more particularly,
to a material used as a heat-resistant photosensitive
material in surface protecting film or interlayer
dielectric for semiconductor device.

BACKGROUND ART

In the semiconductor industries, conventionally,
inorganic materials have been used in interlayer dielectric
materials. However, in recent years, organic materials
having excellent heat resistance, such as polyimide resins,
are being used taking advantage of their properties in the
interlayer dielectric materials.

In forming circuit patterns on semiconductor
integrated circuits or printed circuit boards, the pattern
formation is achieved through a number of cumbersome steps
including formation of a resist layer on the surface of a
base material, exposure of predetermined positions, removal
of unnecessary portions by etching or the like, and
cleaning of the surface of a substrate. Therefore, the
development of heat-resistant photosensitive materials that
constitute resists at necessary portions which can serve as
an insulating material after forming patterns by exposure
and development is desired. For meeting such demands,
recently, photosensitive polyimide and polybenzoxazole are
being mainly used as a resin constituting the resist
wherein the resins themselves have photosensitivity and

hence can easily form a pattern merely by application, exposure, and development, thus shortening the semiconductor fabrication process. Most of the conventional photosensitive polyimide is of a negative type such that an organic solvent is used as a developer and the exposed portions become insoluble. With respect to the negative photosensitive polyimide which creates contrast by a photo-crosslinking reaction, for example, a method has been proposed in which a compound having a photosensitive group is added to or mixed with an acid functional group of a polyimide precursor (see Japanese Patent Application Laid-open No. S54-109828 A).

On the other hand, demands on alkaline aqueous solution development are increasing recently for lowering the material cost and protecting the environment. As an example of conventional techniques, the use of a naphthoquinonediazide compound as a sensitizer and a polybenzoxazole precursor having an acid functional group as a base resin has been proposed (see Japanese Patent Application Publication No. H1-46862).

In the photosensitive resins, the improvement of development properties is important, and therefore the resin is comprised of a polymer having a low molecular weight for increasing the solubility. However, the polymer having a low molecular weight exhibits only unsatisfactory mechanical properties even after being cured, and thus the conventional photosensitive resin materials have a problem in that they cannot exhibit their advantageous resin properties.

The present invention has been achieved in view of the above problem, and an object of the invention is to provide a resin composition comprising a photosensitive polyimide precursor, polybenzoxazole precursor, polyimide, or

polybenzoxazole capable of forming a cured resin having excellent properties, a copolymer thereof, or a mixture thereof; a method for forming a pattern using the resin composition; and an electronic part having an electronic
5 device having the pattern.

DISCLOSURE OF INVENTION

With this situation in mind, the present inventors have conducted intensive studies. As a result, it has been
10 found that, by using a heat-resistant polymer or a precursor thereof having an acid functional group or a substituent derived therefrom and a compound having a substituent derived from an amine functional group in combination, the resin component being cured is presumed to
15 increase in molecular weight, so that a cured resin having desired properties can be formed without sacrificing the development properties.

In the present invention, there are provided a resin composition comprising a photosensitive polyimide
20 precursor, polybenzoxazole precursor, polyimide, or polybenzoxazole capable of forming a cured resin having excellent properties, a copolymer thereof, or a mixture thereof; a method for forming a pattern using the resin composition; and an electronic part having an electronic
25 device having the pattern.

Namely, the present invention is as follows:

[1] A photosensitive resin composition comprising: (A) a polymer having an acid functional group and/or a substituent derived therefrom; (B) a compound having at
30 least one substituent derived from an amine functional group; (C) a photoreactive compound; and (D) a solvent.

[2] The photosensitive resin composition according to [1], wherein the compound of the component (B) further has at

least one acid functional group and/or substituent derived therefrom.

[3] The photosensitive resin composition according to [1], wherein the compound of the component (B) has one

5 substituent or two substituents derived from an amine functional group.

[4] The photosensitive resin composition according to [3], wherein the component (B) has one or two acid functional group and/or substituent derived therefrom.

10 [5] The photosensitive resin composition according to [1], wherein the component (A) further has at least one substituent derived from an amine functional group in itself or in another component (A), and wherein the component (B) further has at least one acid functional

15 group and/or substituent derived therefrom.

[6] The photosensitive resin composition according to [5], wherein the component (A) has one substituent derived from an amine functional group, and wherein the component (B) has one substituent or two substituents derived from an

20 amine functional group and has one acid functional group and/or substituent derived therefrom.

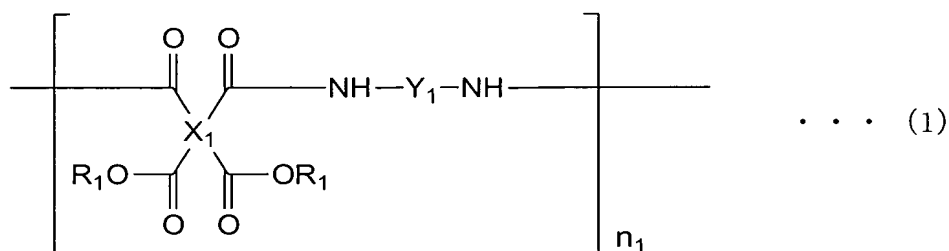
[7] The photosensitive resin composition according to [1], wherein the polymer (A) is a heat-resistant polymer.

[8] The photosensitive resin composition according to [1],

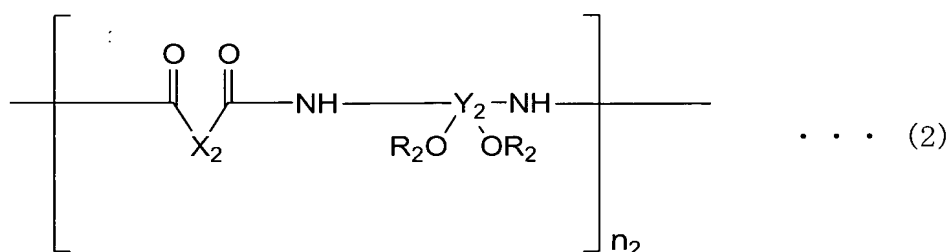
25 wherein the acid functional group in the polymer (A) is a carboxyl group and/or a phenolic hydroxyl group.

[9] The photosensitive resin composition according to [7] or [8], wherein the heat-resistant polymer is a polyimide precursor represented by the general formula (1) below or

30 polyimide derived from the precursor, a polybenzoxazole precursor represented by the general formula (2) below or polybenzoxazole derived from the precursor, a copolymer thereof, or a mixture thereof:



wherein X_1 represents a tetravalent organic group, Y_1
 5 represents a divalent organic group, R_1 represents hydrogen
 or a monovalent organic group, and n_1 represents an integer
 of 2 to 500 corresponding to the number of the repeating
 units of the polymer,



10

wherein X_2 represents a divalent organic group, Y_2
 represents a tetravalent organic group, of which two
 valences are used in bonding to hydroxyl groups, R_2
 15 represents hydrogen or a monovalent organic group, and n_2
 represents an integer of 2 to 500 corresponding to the
 number of the repeating units of the polymer.

[10] A method for forming a pattern, comprising: a step of
 applying the photosensitive resin composition according to
 20 [1] onto a substrate and drying it; a step of subjecting
 the applied and dried photosensitive resin film to light
 exposure; a step of subjecting the exposed photosensitive
 resin film to development using an alkaline aqueous

solution; and a step of subjecting the pattern obtained in the development to heat treatment.

[11] An electronic part having an electronic device having the pattern obtained by the method according to [10],

5 wherein the pattern is an interlayer dielectric layer and/or a surface protecting film layer.

The invention according to any one of the items [1] to [8] is directed to a photosensitive resin composition which comprises: (A) a polymer having an acid functional group
10 and/or a substituent derived therefrom; (B) a compound having at least one substituent derived from an amine functional group; (C) a photoreactive compound; and (D) a solvent, and which forms, utilizing an acid functional group or a substituent derived therefrom of a conventional
15 heat-resistant polymer or a precursor thereof, a cured resin having desired properties from the heat-resistant polymer or precursor thereof without sacrificing the development properties.

The invention according to the item [7] has an effect
20 of the invention according to the item [1] and forms a cured resin having more excellent properties, and the invention according to the item [8] is directed to the photosensitive resin composition wherein the acid functional group in polymer (A) is a carboxyl group and/or
25 a phenolic hydroxyl group.

The invention according to the item [9] is directed to the photosensitive resin composition wherein polymer (A) is a polyimide precursor represented by the general formula (1), a polybenzoxazole precursor represented by the general
30 formula (2), polyimide or polybenzoxazole derived from the precursor, a copolymer thereof, or a mixture thereof, and is directed to the resin composition having an effect of the invention according to the item [1], wherein polymer

(A) is a photosensitive polyimide precursor, a polybenzoxazole precursor, polyimide, polybenzoxazole, a copolymer thereof, or a mixture thereof, which forms a cured resin having more excellent properties.

5 The invention according to the item [10] is directed to a method for forming a pattern using the heat-resistant photosensitive resin composition having an effect of the invention according to the item [1]. The invention according to the item [11] is directed to an electronic
10 part having an electronic device with the pattern obtained by the method according to the item [10] as a surface protecting film layer and/or an interlayer dielectric layer.

15 BRIEF DESCRIPTION OF DRAWING

Fig. 1 is a manufacturing process diagram of a semiconductor device of a multilayer wiring structure.

BEST MODE FOR CARRYING OUT THE INVENTION

20 Embodiments of the present invention will be explained below in detail with reference to the accompanying drawing. Note that the invention is not limited to the embodiments.

In the present invention, the essential components include (A) a polymer having an acid functional group
25 and/or a substituent derived therefrom, (B) a compound having at least one substituent derived from an amine functional group, (C) a photoreactive compound, and (D) a solvent. It is considered that component (B) is reacted with the acid functional group and/or substituent derived
30 therefrom in component (A) due to heating to cause polymer (A) to have a higher molecular weight.

With respect to component (A) in the present invention, examples of the polymers having an acid

functional group and/or a substituent derived therefrom may include polyimide precursors obtained from a tetracarboxylic dianhydride and a diamine, polybenzoxazole precursors obtained from a dicarboxylic acid and a bisaminophenol, copolymers thereof, and mixtures thereof. The polymer has in its molecule an acid functional group, such as a carboxylic acid group or an aromatic hydroxyl group, and/or a substituent derived therefrom, and can further have a substituent derived from an amine functional group. It is considered that the molecular weight of the polymer of component (A) can be increased utilizing the functional group.

For example, the compound of component (B) having at least one substituent derived from an amine functional group is reacted with the polymer of component (A) to increase the molecular weight of the polymer. It is considered that, for example, the compound having at least one substituent derived from an amine functional group and being bonded to the polymer is further reacted with the polymer or a polymer of another component (A), that is, the compound forms crosslinking in the polymer molecules or between the polymer molecules to further increase the molecular weight of the polymer of component (A). With respect to the manner for increasing the molecular weight of the polymer of component (A), there is no particular limitation.

Examples of tetracarboxylic dianhydrides may include pyromellitic dianhydride, 3,3',4,4'-biphenyltetracarboxylic dianhydride, 2,3,3',4'-biphenyltetracarboxylic dianhydride, 2,2',3,3'-biphenyltetracarboxylic dianhydride, 3,3',4,4'-benzophenonetetracarboxylic dianhydride, 2,2',3,3'-benzophenonetetracarboxylic dianhydride, 2,2-bis(3,4-dicarboxyphenyl)propane dianhydride, 2,2-bis(2,3-

dicarboxyphenyl)propane dianhydride, 1,1-bis(3,4-dicarboxyphenyl)ethane dianhydride, 1,1-bis(2,3-dicarboxyphenyl)ethane dianhydride, bis(3,4-dicarboxyphenyl)methane dianhydride, bis(2,3-dicarboxyphenyl)methane dianhydride, bis(3,4-dicarboxyphenyl) sulfone dianhydride, bis(3,4-dicarboxyphenyl) ether dianhydride, 1,2,3,4-cyclopentanetetracarboxylic dianhydride, 2,2-bis[4-(4-aminophenoxy)phenyl]propane, 1,2,5,6-naphthalenetetracarboxylic dianhydride, 2,3,6,7-naphthalenetetracarboxylic dianhydride, 2,3,5,6-pyridinetetracarboxylic dianhydride, 3,4,9,10-perylenetetracarboxylic dianhydride, 3,3',4,4'-tetraphenylsilanetetracarboxylic dianhydride, 2,2-bis(3,4-dicarboxyphenyl)hexafluoropropane dianhydride, N-(trimellitic dianhydride)-2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, N-(trimellitic dianhydride)-2,2'-bis(3-amino-4-hydroxyphenyl)propane, N-(trimellitic dianhydride)-4,4'-diamino-3,3'-dihydroxybiphenyl, N-(trimellitic dianhydride)-2,2'-bis(3-amino-4-hydroxyphenyl) sulfone, N-(trimellitic dianhydride)-2,2'-bis(3-amino-4-hydroxyphenyl) ether, N-(trimellitic dianhydride)-2,2'-bis(3-amino-4-hydroxyphenyl)propane, N-(trimellitic dianhydride)-2,4-diaminophenol, N-(trimellitic dianhydride)-2,5-diaminophenol, and N-(trimellitic dianhydride)-1,4-diamino-2,5-dihydroxybenzene, and these may be used individually or in combination. The tetracarboxylic dianhydride is not necessarily limited to those listed above. The tetracarboxylic dianhydride and the diamine may be used in combination.

Examples of diamines may include 4,4'-diaminodiphenyl ether, 4,4'-diaminodiphenylmethane, 4,4'-diaminodiphenyl

sulfone, 4,4'-diaminodiphenyl sulfide, m-phenylenediamine, p-phenylenediamine, 1,5-naphthalenediamine, 2,6-naphthalenediamine, bis(4-aminophenoxyphenyl) sulfone, bis(3-aminophenoxyphenyl) sulfone, bis(4-aminophenoxy)biphenyl, bis[4-(4-aminophenoxy)phenyl] ether, 1,4-bis(4-aminophenoxy)benzene, 2,2'-dimethyl-4,4'-diaminobiphenyl, 2,2'-diethyl-4,4'-diaminobiphenyl, 3,3'-dimethyl-4,4'-diaminobiphenyl, 3,3'-diethyl-4,4'-diaminobiphenyl, 2,2',3,3'-tetramethyl-4,4'-diaminobiphenyl, 2,2',3,3'-tetraethyl-4,4'-diaminobiphenyl, 2,2'-dimethoxy-4,4'-diaminobiphenyl, 3,3'-dimethoxy-4,4'-diaminobiphenyl, 2,2'-dihydroxy-4,4'-diaminobiphenyl, 3,3'-dihydroxy-4,4'-diaminobiphenyl, 2,2'-di(trifluoromethyl)-4,4'-diaminobiphenyl, 2,4-diaminobenzoic acid, 3,5-diaminobenzoic acid, 3,3'-diaminobiphenyl-5,5'-dicarboxylic acid, 4,4'-diaminodiphenyl ether-5,5'-dicarboxylic acid, 4,4'-diaminodiphenylmethane-5,5'-dicarboxylic acid, 4,4'-diaminodiphenyl sulfone-5,5'-dicarboxylic acid, 4,4'-diaminodiphenyl sulfide-5,5'-dicarboxylic acid, 4,4'-diamino-3,3'-dihydroxybiphenyl, 2,2'-bis(3-amino-4-hydroxyphenyl)propane, 2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, oxybis(3-amino-4-hydroxyphenyl), bis(3-amino-4-hydroxyphenyl) sulfone, 2,4-diaminophenol, 1,4-diamino-2,5-dihydroxybenzene, N,N'-(4-aminophenylcarbonyl)-3,3'-dihydroxybiphenyl, N,N'-(3-aminophenylcarbonyl)-3,3'-dihydroxybiphenyl, N,N'-(4-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, N,N'-(3-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, N,N'-(4-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)propane, N,N'-(3-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)propane, N,N'-(4-

aminophenylcarbonyl)-oxybis(3-amino-4-hydroxyphenyl), N,N'-(3-aminophenylcarbonyl)-oxybis(3-amino-4-hydroxyphenyl), N,N'-(4-aminophenylcarbonyl)-bis(3-amino-4-hydroxyphenyl) sulfone, N,N'-(3-aminophenylcarbonyl)-bis(3-amino-4-hydroxyphenyl) sulfone, N,N'-(4-aminophenylcarbonyl)-2,4-diaminophenol, N,N'-(3-aminophenylcarbonyl)-2,4-diaminophenol, N,N'-(4-aminophenylcarbonyl)-1,4-diamino-2,5-dihydroxybenzene, and N,N'-(3-aminophenylcarbonyl)-1,4-diamino-2,5-dihydroxybenzene, and these may be used individually or in combination. The diamine is not necessarily limited to those listed above.

As mentioned above, a tetracarboxylic dianhydride is reacted with a diamine to obtain a polyamide acid as a polyimide precursor, and the introduction of the functional group R₁ in the general formula (1) enables control of the solubility during the development and/or patterning using a photoreaction.

When imparting negative photosensitivity to a resin composition, it is preferred that the functional group contains a reactive functional group, such as a vinyl group, an allyl group, an acryloyl group, or a methacryloyl group from the viewpoint of obtaining an excellent pattern form after the development.

As examples of the reactive functional groups, there can be mentioned functional groups having a carbon-carbon unsaturated double bond. Examples of functional groups having a carbon-carbon unsaturated double bond may include allyl alcohol, 2-methyl-2-propen-1-ol, crotyl alcohol, 3-buten-1-ol, 3-buten-2-ol, 3-methyl-2-buten-1-ol, 2-methyl-3-buten-1-ol, 3-methyl-3-buten-1-ol, 2-methyl-3-buten-2-ol, 2-penten-1-ol, 4-penten-1-ol, 3-penten-2-ol, 4-penten-2-ol, 1-penten-3-ol, 4-methyl-3-penten-1-ol, 3-methyl-1-penten-3-ol, 2-hexen-1-ol, 3-hexen-1-ol, 4-hexen-1-ol, 5-hexen-1-ol,

1-hexen-3-ol, 1-heptan-3-ol, 6-methyl-5-heptan-2-ol, 1-octan-3-ol, citronellol, 3-nonen-1-ol, 5-decan-1-ol, 9-decan-1-ol, 7-decan-1-ol, 1,4-pentadien-3-ol, 2,4-hexadien-1-ol, 1,5-hexadien-3-ol, 1,6-heptadien-4-ol, 2,4-dimethyl-
5 2,6-heptadien-1-ol, nerol, geraniol, linalool, 2-cyclohexen-1-ol, 3-cyclohexen-1-methanol, isopulegol, 5-norbornen-2-ol, 5-norbornen-2-methanol, ethylene glycol vinyl ether, 1,4-butanediol vinyl ether, 1,6-hexanediol vinyl ether, diethylene glycol vinyl ether, 2-hydroxyethyl
10 acrylate, 3-hydroxypropyl acrylate, 2-hydroxyethyl methacrylate, 3-hydroxypropyl methacrylate, 4-hydroxybutyl acrylate, 4-hydroxybutyl methacrylate, pentaerythritol diacrylate monostearate, pentaerythritol triacrylate, caprolactone 2-(methacryloyloxy)ethyl ester, dicaprolactone
15 2-(methacryloyloxy)ethyl ester, 2-hydroxy-3-phenoxypropyl acrylate, and 2-hydroxy-3-phenoxypropyl methacrylate. The functional group having a carbon-carbon unsaturated double bond is not necessarily limited to those listed above. The functional group may be partially introduced to impart
20 alkali solubility to the polymer. When the polymer has an acid functional group at its end, the functional group can be introduced.

When imparting positive photosensitivity to a resin composition, a substituent may be introduced to control the
25 solubility or utilize an elimination reaction. Examples of such substituents may include methyl, ethyl, propyl, isopropyl, n-butyl, s-butyl, t-butyl, cyclopropenyl, cyclobutyl, cyclopentyl, cyclohexyl, cyclohexylmethyl, cyclohexenyl, norbornyl, norbornenyl, adamantyl, benzyl, p-nitrobenzyl, trifluoromethyl, methoxyethyl, ethoxyethyl,
30 methoxymethyl, ethoxymethyl, methoxyethoxymethyl, benzoxyethyl, ethoxytetrahydropyranyl, tetrahydrofuranyl, 2-trimethylsilylethoxymethyl, trimethylsilyl, t-

butyldimethylsilyl, 3-oxocyclohexyl, 9-fluorenylmethyl, and methylthiomethyl. The substituent is not necessarily limited to those listed above. The functional group may be partially introduced to control the alkali solubility.

5 When the polymer has an acid functional group at its end, the functional group can be introduced.

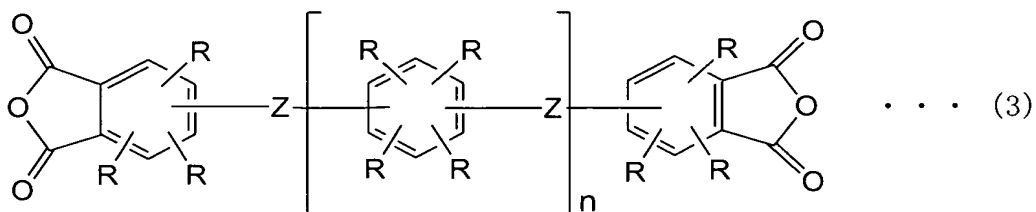
Examples of dicarboxylic acids may include 3-fluoroisophthalic acid, 2-fluoroisophthalic acid, 3-fluorophthalic acid, 2-fluorophthalic acid, 2,4,5,6-
10 tetrafluoroisophthalic acid, 3,4,5,6-tetrafluorophthalic acid, 4,4'-hexafluoroisopropylidenediphenyl-1,1'-dicarboxylic acid, perfluorosuberic acid, 2,2'-bis(trifluoromethyl)-4,4'-biphenylenedicarboxylic acid, terephthalic acid, isophthalic acid, and 4,4'-oxydiphenyl-
15 1,1'-dicarboxylic acid, and these may be used individually or in combination. The dicarboxylic acid is not necessarily limited to those listed above.

Examples of bisaminophenols may include 2,2'-dihydroxy-4,4'-diaminobiphenyl, 3,3'-dihydroxy-4,4'-
20 diaminobiphenyl, 4,4'-diamino-3,3'-dihydroxybiphenyl, 2,2'-bis(3-amino-4-hydroxyphenyl)propane, 2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, oxybis(3-amino-4-hydroxyphenyl), bis(3-amino-4-hydroxyphenyl) sulfone, 2,4-diaminophenol, 1,4-diamino-2,5-dihydroxybenzene, N,N'-(4-aminophenylcarbonyl)-3,3'-dihydroxybiphenyl, N,N'-(3-aminophenylcarbonyl)-3,3'-dihydroxybiphenyl, N,N'-(4-aminophenylcarbonyl)
25 2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, N,N'-(3-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)hexafluoropropane, N,N'-(4-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)propane, N,N'-(3-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)propane, N,N'-(4-aminophenylcarbonyl)2,2'-bis(3-amino-4-hydroxyphenyl)propane, N,N'-(4-

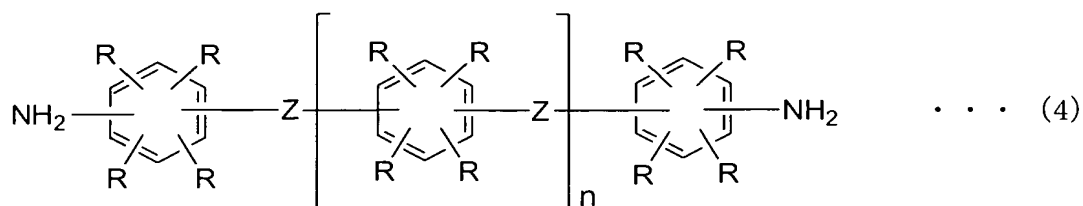
aminophenylcarbonyl)-oxybis(3-amino-4-hydroxyphenyl), N,N'-(3-aminophenylcarbonyl)-oxybis(3-amino-4-hydroxyphenyl), N,N'-(4-aminophenylcarbonyl)-bis(3-amino-4-hydroxyphenyl) sulfone, N,N'-(3-aminophenylcarbonyl)-bis(3-amino-4-hydroxyphenyl) sulfone, N,N'-(4-aminophenylcarbonyl)-2,4-diaminophenol, N,N'-(3-aminophenylcarbonyl)-2,4-diaminophenol, N,N'-(4-aminophenylcarbonyl)-1,4-diamino-2,5-dihydroxybenzene, and N,N'-(3-aminophenylcarbonyl)-1,4-diamino-2,5-dihydroxybenzene, and these may be used individually or in combination. The bisaminophenol is not necessarily limited to those listed above. The bisaminophenol and the above diamine may be used in combination.

Like in the polyamide acid as the polyimide precursor, in the polybenzoxazole precursor, the introduction of a functional group similar to R₁ in the general formula (1) into R₂ in the general formula (2) enables control of the solubility during the development and/or patterning using a photoreaction. The functional group may be partially introduced to control the alkali solubility. When the polymer has an acid functional group at its end, the functional group can be introduced.

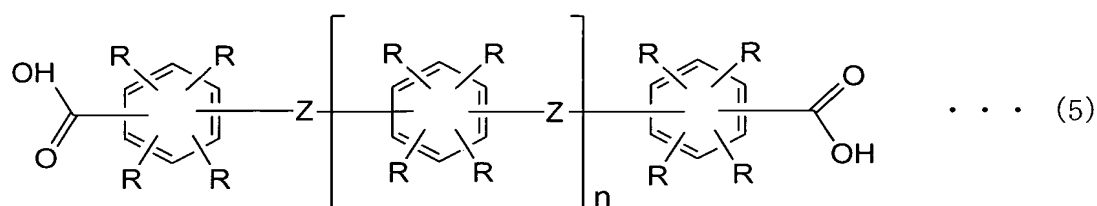
More generally, the tetracarboxylic dianhydride is represented by the following general formula (3).



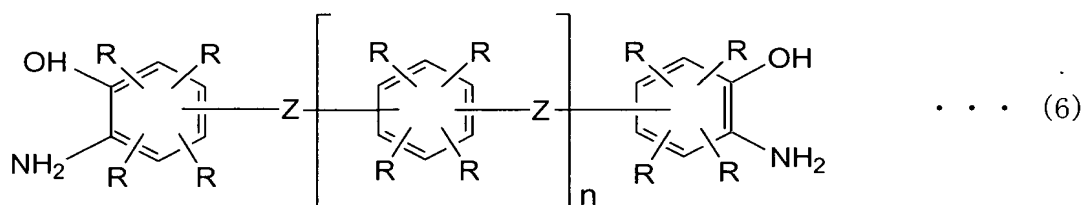
More generally, the diamine is represented by the following general formula (4).



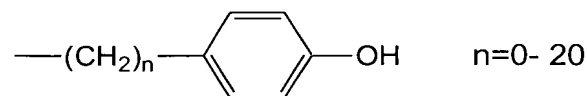
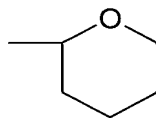
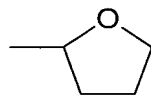
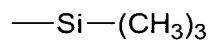
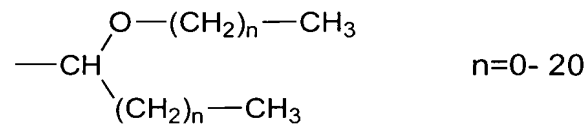
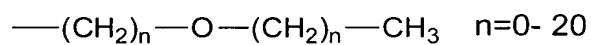
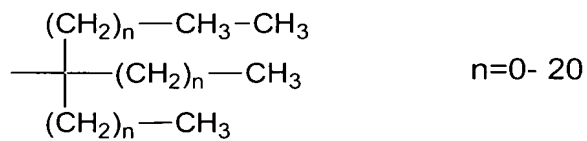
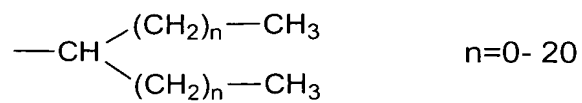
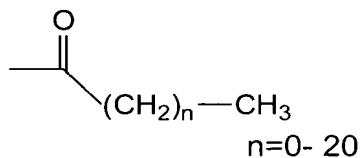
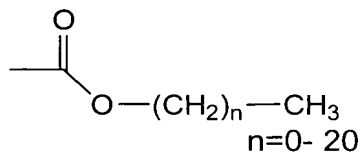
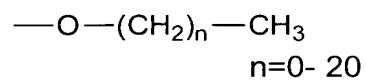
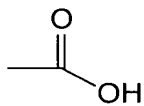
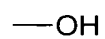
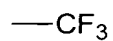
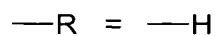
More generally, the dicarboxylic acid is represented
5 by the following general formula (5).

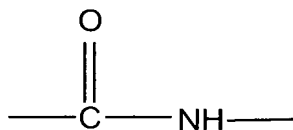
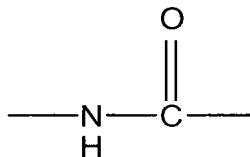
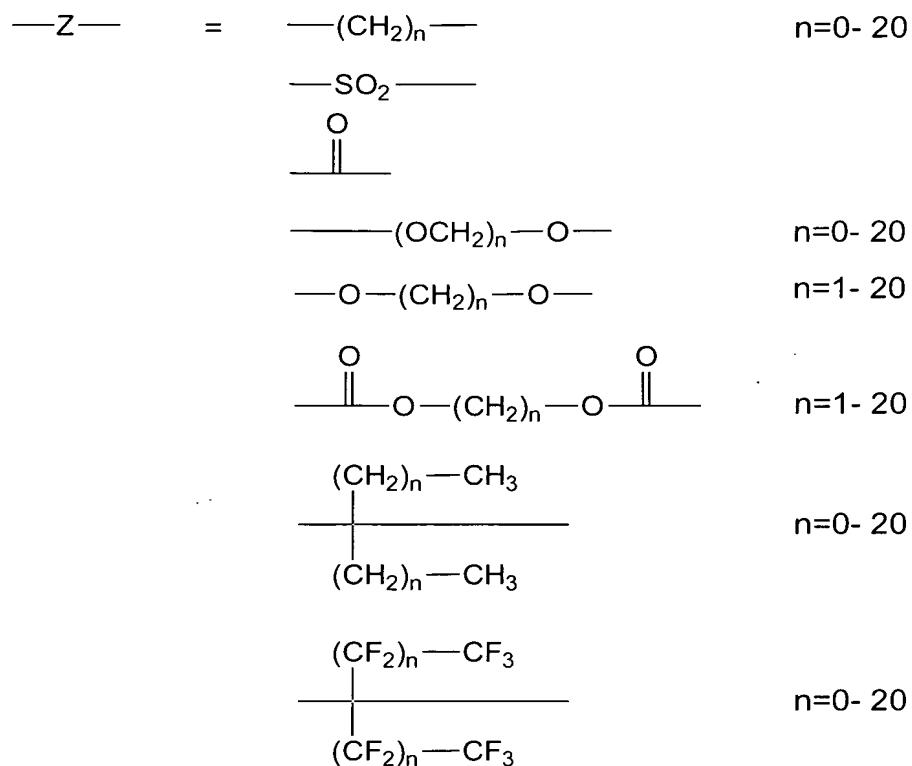


More generally, the bisaminophenol is represented by
10 the following general formula (6).



In the general formulae (3) to (6), n is 0 or a
15 natural number, and characters R and Z are a monovalent organic group and a divalent organic group, respectively, and represented by the following formulae.





- 5 It is preferred that polymer (A) has at its end at least one substituent derived from an amine functional group and/or at least one acid functional group and/or substituent derived therefrom from the viewpoint of forming a cured resin having further more excellent properties.
- 10 When the amine functional group is a primary amine, the stability of the photosensitive resin composition is lowered due to a side reaction, and therefore it is

preferred that at least one of the two hydrogen atoms on the amine functional group is replaced by another atom or another functional group from the viewpoint of obtaining the photosensitive resin composition having satisfactory stability.

Examples of the substituents on nitrogen derived from an amine functional group may include carbonyl, carbamate, amide, sulfonyl, sulfenyl, phosphinyl, and alkylsilyl. Of these, preferred are carbonyl, carbamate, and sulfonyl from the viewpoint of obtaining a cured resin having more excellent properties.

The substituent is hydrogen or any of those represented by the following general formulae (7) to (9):



15

wherein R_3 represents a monovalent organic group, preferably having 1 to 20 carbon atoms; X_3 represents an oxygen, sulfur, or nitrogen atom; and n represents 1 when X_3 is an oxygen atom or a sulfur atom, or n represents 2 when X_3 is a nitrogen atom.

Examples of R_3 's may include methyl, ethyl, propyl, isopropyl, n-butyl, s-butyl, t-butyl, cyclopropenyl, cyclobutyl, cyclopentyl, cyclohexyl, cyclohexylmethyl, cyclohexenyl, norbornyl, norbornenyl, adamantyl, benzyl, p-nitrobenzyl, trifluoromethyl, methoxyethyl, ethoxyethyl,

25

methoxymethyl, ethoxymethyl, methoxyethoxymethyl,
benzoxymethyl, ethoxytetrahydropyranyl, tetrahydrofuranyl,
2-trimethylsilylethoxymethyl, trimethylsilyl, t-
butyldimethylsilyl, 3-oxocyclohexyl, 9-fluorenylmethyl,
5 methylthiomethyl, phenyl, toluyl, xylyl, 9,10-
dihydroanthranyl, trimethylphenyl, pentamethylphenyl,
biphenyl, terphenyl, quaterphenyl, dimethylbiphenyl,
naphthalenyl, methylnaphthalenyl, fluorenyl, fluorophenyl,
fluorobiphenyl, isopropylidenebiphenyl,
10 tetrafluoroisopropylidenebiphenyl, benzyl phenyl ether,
phenyl ether, phenoxytoluoyl, methoxybiphenyl,
dimethoxybiphenyl, methoxynaphthalenyl,
dimethoxynaphthalenyl, and nitrophenyl. R₃ is not
necessarily limited to those listed above.

15 As the acid functional group and/or substituent
derived therefrom, one represented by R₁ in the general
formula (1) or R₂ in the general formula (2)~~(1)~~ may be
used.

In the present invention, basically it is preferred
20 that compound (B) is a compound having at least one
substituent derived from an amine functional group. When
the amine functional group is a primary amine, the
stability of the photosensitive resin composition may be
lowered due to a side reaction. Therefore at least one of
25 the two hydrogen atoms on the primary amine functional
group is to be replaced by another atom or another
functional group.

More generally, the compound is represented by the
following general formula (10):

30



wherein X_4 represents an n-valent organic group, each of R_4 and R_5 represents hydrogen or a monovalent organic group, and n represents an integer of 2 to 6.

Examples of X_4 's may include aliphatic groups, such as
5 an alkyl chain, a cyclopentyl ring, a cyclohexyl ring, a cyclooctyl ring, and a bicyclo ring, and these may contain a hetero atom, such as oxygen or sulfur.

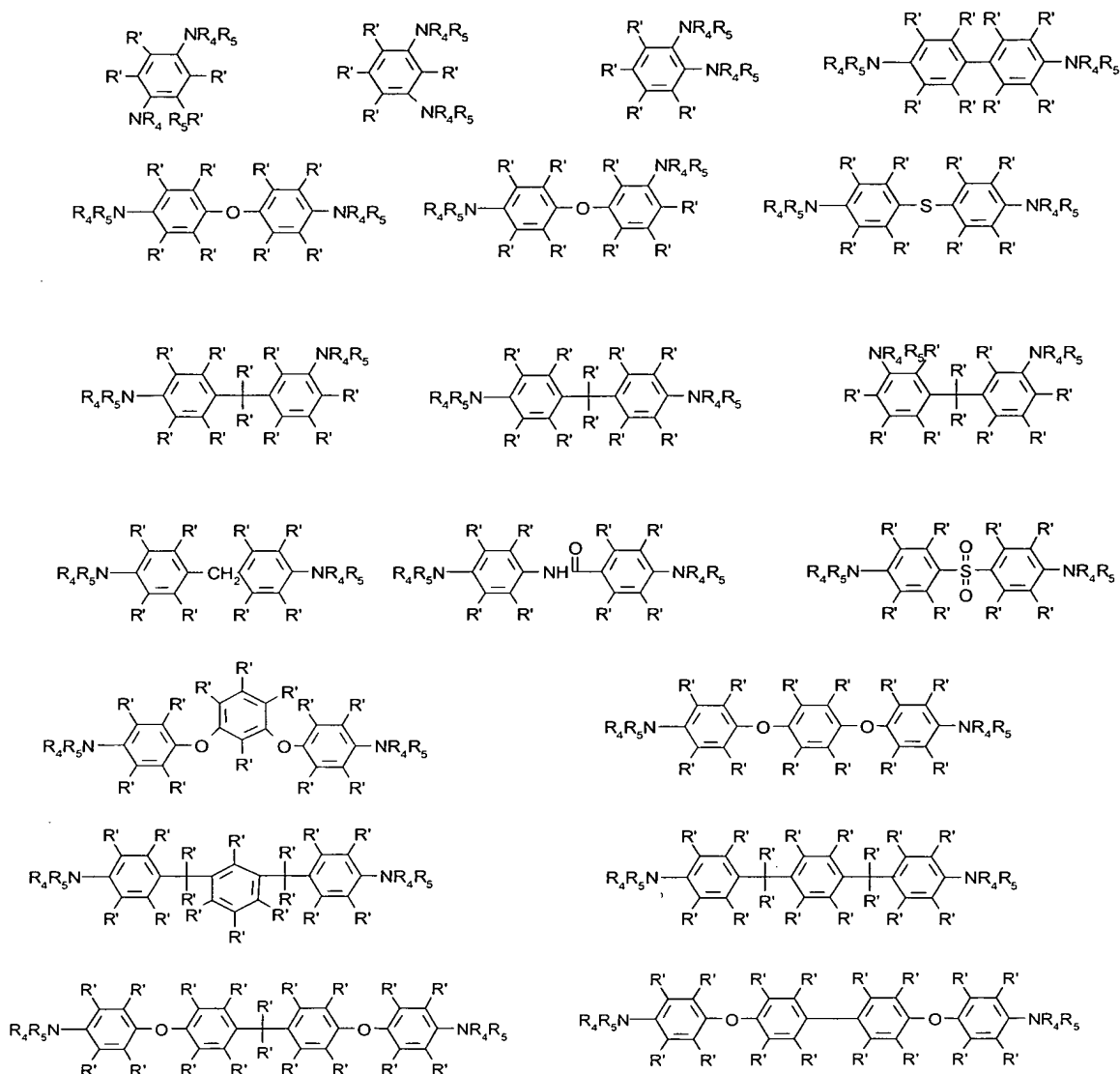
Further, examples of X_4 's may include aromatic groups, such as a benzene ring and a naphthalene ring. Of these,
10 preferred are aromatic groups from the viewpoint of achieving excellent heat resistance. Examples of aromatic groups may include benzene, toluene, cumene, diphenylmethane, xylene, 9,10-dihydroanthracene, mesitylene, hexamethylbenzene, biphenyl, terphenyl,
15 triphenylbenzene, quaterphenyl, dimethylbiphenyl, azulene, naphthalene, methylnaphthalene, anthracene, fluorene, fluorobenzene, fluorobiphenyl, anisole, benzyl phenyl ether, phenyl ether, phenoxytoluene, tolyl ether, methoxybiphenyl, dimethoxybiphenyl, methoxynaphthalene,
20 dimethoxynaphthalene, nitrobenzene, acetophenone, benzophenone, benzanilide, phenyl sulfone, furan, and thiophene. X_4 is not necessarily limited to those listed above.

Each of R_4 and R_5 in general formula (10) is generally
25 hydrogen or any of those represented by the general formula (11) below. Examples of R_4 's or R_5 's may include carbonyl, carbamate, amide, sulfonyl, sulfenyl, phosphinyl, and alkylsilyl. Of these, preferred are carbonyl, carbamate, and sulfonyl from the viewpoint of obtaining a cured resin
30 having more excellent properties. As the substituent, hydrogen or one represented by the general formulae (7) to (9) may be used.



wherein X_4 represents an $(m+n+1)$ -valent organic group; each
of R_4 , R_5 , R_6 , and R_7 represents hydrogen or a monovalent
5 organic group; n represents 1 to 6; and each of l and m
represents 1 to 4, wherein each of R_6 and R_7 is a
monovalent organic group and a functional group similar to
 R_1 or R_2 above may be introduced.

Structures of more preferred compounds of the general
10 formula (10) are shown below.



In the aforementioned formulae, R' represents hydrogen or a monovalent organic group, and examples of R' 's may include R_3 in the general formulae (7) to (9), an ether

5 group, an ester group, and a halogen, such as fluorine.

The compound represented by the general formula (10) may be a compound having at least one substituent derived from an amine functional group and further having at least

10 one acid functional group and/or substituent derived therefrom. The acid functional group is more preferably a carboxylic acid group and/or a phenolic hydroxyl group.

Specifically, the compound is represented by the following general formula (11):



5

wherein X_4 represents an $(m+n+1)$ -valent organic group; each of R_4 , R_5 , R_6 , and R_7 represents hydrogen or a monovalent organic group; n represents 1 to 6; and each of l and m represents 1 to 4, wherein each of R_6 and R_7 is a
10 monovalent organic group and a functional group similar to R_1 or R_2 above may be introduced.

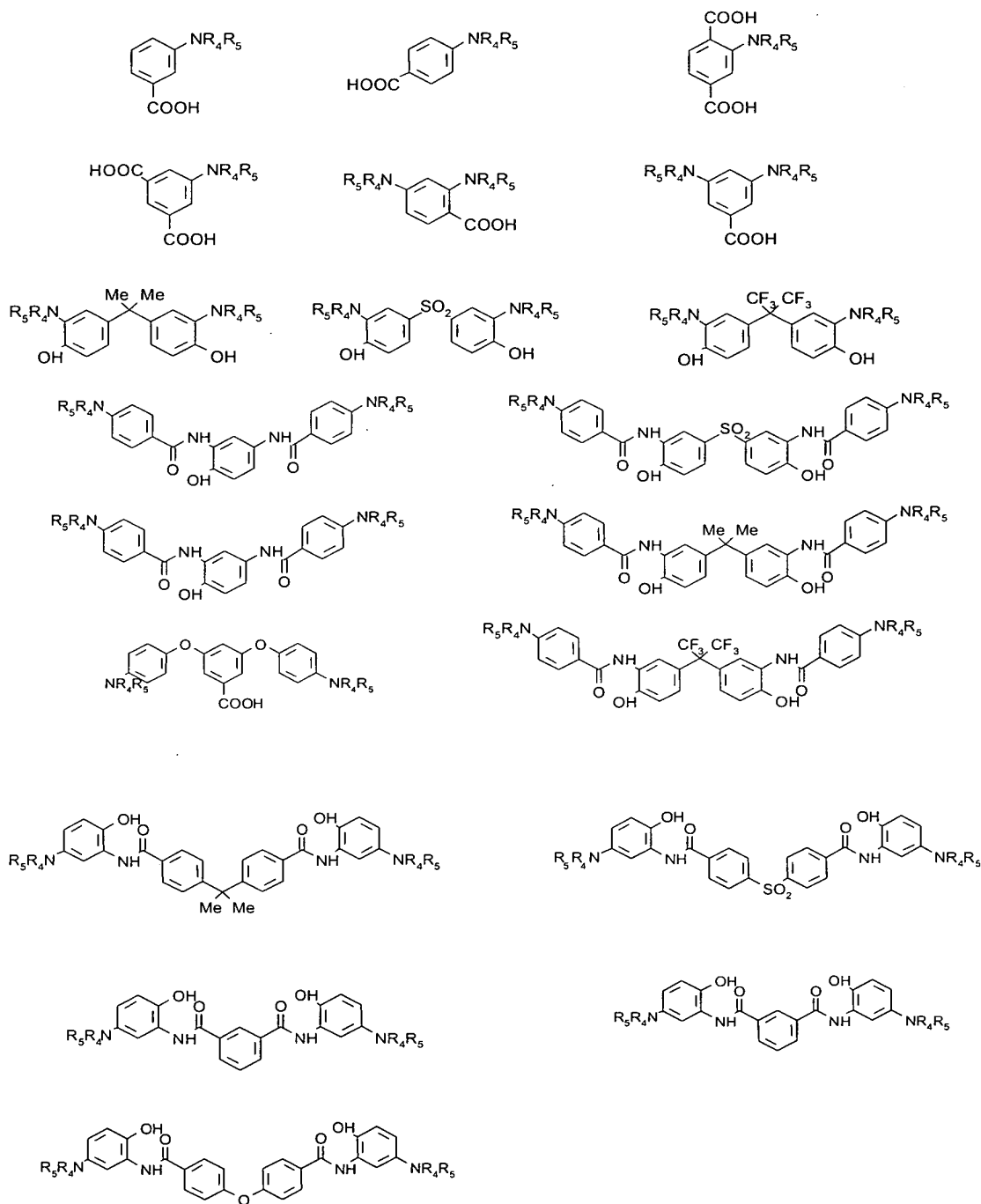
Examples of X_4 's may include aliphatic groups, such as an alkyl chain, a cyclopentyl ring, a cyclohexyl ring, a cyclooctyl ring, and a bicyclo ring, and these may contain
15 a hetero atom, such as oxygen or sulfur.

Further, examples of X_4 's may also include aromatic groups, such as a benzene ring and a naphthalene ring. Of these, preferred are aromatic groups from the viewpoint of achieving excellent heat resistance.

20 Examples of aromatic groups may include benzene, toluene, cumene, diphenylmethane, xylene, 9,10-dihydroanthracene, mesitylene, hexamethylbenzene, biphenyl, terphenyl, triphenylbenzene, quaterphenyl, dimethylbiphenyl, azulene, naphthalene, methylnaphthalene,
25 anthracene, fluorene, fluorobenzene, fluorobiphenyl, anisole, benzyl phenyl ether, phenyl ether, phenoxytoluene, tolyl ether, methoxybiphenyl, dimethoxybiphenyl, methoxynaphthalene, dimethoxynaphthalene, nitrobenzene, acetophenone, benzophenone, benzanilide, phenyl sulfone,
30 furan, and thiophene. X_4 is not necessarily limited to those listed above.

Each of R_4 and R_5 in the general formula (11) is generally hydrogen or any of those represented by the general formulae (7) to (9). Examples of R_4 's or R_5 's may include carbonyl, carbamate, amide, sulfonyl, sulfenyl, phosphinyl, and alkylsilyl. Of these, preferred are carbonyl, carbamate, and sulfonyl from the viewpoint of obtaining a cured resin having more excellent properties. As the substituent, hydrogen or one represented by the general formulae (7) to (9) may be used.

Structures of more preferred compounds of the general formula (11) are shown below.



Hydrogen on the aromatic ring may be replaced by a monovalent organic group, such as R_3 in the general formulae (7) to (9), or a halogen, such as fluorine.

It is considered that compound (B) serves as a chain extender capable of increasing the molecular weight of

polymer (A) during the heat curing. Therefore, the compound is preferably incorporated in an amount in the range of from 0.05 to 50 parts by weight, more preferably 0.2 to 20 parts by weight, relative to 100 parts by weight of polymer component (A).

Photoreactive compound (C) used in the present invention is a sensitizer, and, when the compound is a negative sensitizer, it has photopolymerizability and reduces the solubility of the exposed portion.

Examples of the sensitizers may include benzophenones, such as benzophenone, Michler's ketone, 4,4,-bis(diethylamino)benzophenone, and 3,3,4,4,-tetra(t-butylperoxycarbonyl)benzophenone; benzylidenes, such as 3,5-bis(diethylaminobenzylidene)-N-methyl-4-piperidone and 3,5-bis(diethylaminobenzylidene)-N-ethyl-4-piperidone; coumarins, such as 7-diethylamino-3-thenoylcoumarin, 4,6-dimethyl-3-ethylaminocoumarin, 3,3,-carbonylbis(7-diethylaminocoumarin), 7-diethylamino-3-(1-methylbenzimidazolyl) coumarin ~~7-diethylamino-3-(1-methylmethybenzimidazolyl) coumarin~~, and 3-(2-benzothiazolyl)-7-diethylaminocoumarin; anthraquinones, such as 2-t-butylanthraquinone, 2-ethylanthraquinone, and 1,2-benzanthraquinone; benzoin ethers, such as benzoin methyl ether, benzoin ethyl ether, and benzoin isopropyl ether; thioxanthenes, such as 2,4-dimethylthioxanthone, 2,4-diethylthioxanthone, 2,4-diisopropylthioxanthone, and 2-isopropylthioxanthone; mercapto compounds, such as ethylene glycol di(3-mercaptopropionate), 2-mercaptobenzothiazole, 2-mercaptobenzoxazole, and 2-mercaptobenzimidazole; glycines, such as N-phenylglycine, N-methyl-N-phenylglycine, N-ethyl-N-(p-chlorophenyl)glycine, and N-(4-cyanophenyl)glycine; oximes, such as 1-phenyl-1,2-butanedione-2-(o-methoxycarbonyl)oxime, 1-phenyl-1,2-

propanedione-2-(o-methoxycarbonyl)oxime, 1-phenyl-1,2-propanedione-2-(o-ethoxycarbonyl)oxime, and 1-phenyl-1,2-propanedione-2-(o-benzoyl)oxime; and 2,2'-bis(o-chlorophenyl)-4,4',5,5'-tetraphenylbiimidazole. These may
5 be used individually or in combination. The sensitizer is not necessarily limited to those listed above.

These sensitizers may be used individually or in combination. The amount of the sensitizer used is generally in the range of from 0.01 to 15% by weight, or
10 the total amount of the two or more sensitizers used is preferably in the range of from 0.1 to 40% by weight, more preferably 1 to 20 parts by weight, relative to 100 parts by weight of polymer component (A).

A positive sensitizer generates acid and increases the
15 solubility of the exposed portion in an alkaline aqueous solution. Examples of positive sensitizers may include o-quinonediazide compounds, aryldiazonium salts, diaryliodonium salts, and triarylsulfonium salts. The positive sensitizer is not limited to the compounds listed
20 above, and any compound that generates acid as a response to light exposure may be used.

The o-quinonediazide compound may be obtained by subjecting, for example, o-quinonediazidosulfonyl chloride, a hydroxy compound, and an amino compound or the like to
25 condensation reaction in the presence of a dehydrochlorination agent. Examples of the o-quinonediazidosulfonyl chlorides may include benzoquinone-1,2-diazido-4-sulfonyl chloride, naphthoquinone-1,2-diazido-5-sulfonyl chloride, and naphthoquinone-1,2-
30 diazido-4-sulfonyl chloride. The o-quinonediazidosulfonyl chloride is not necessarily limited to those listed above.

As the hydroxy compound, for example, hydroquinone, resorcinol, pyrogallol, bisphenol A, bis(4-

hydroxyphenyl)methane, 2,2'-bis(4-hydroxyphenyl)hexafluoropropane, 2,3,4-trihydroxybenzophenone, 2,3,4,4'-tetrahydroxybenzophenone, 2,2',4,4'-tetrahydroxybenzophenone, 2,3,4,2',3'-pentahydroxybenzophenone, 2,3,4,3',4',5'-hexahydroxybenzophenone, bis(2,3,4-trihydroxyphenyl)methane, bis(2,3,4-trihydroxyphenyl)propane, 4b,5,9b,10-tetrahydro-1,3,6,8-tetrahydroxy-5,10-dimethylindeno[2,1-a]indene, tris(4-hydroxyphenyl)methane, or tris(4-hydroxyphenyl)ethane may be used. The hydroxy compound is not necessarily limited to those listed above.

Examples of the aryldiazonium salts, diaryliodonium salts, or triarylsulfonium salts may include benzenediazonium p-toluenesulfonate, diphenyliodonium 9,10-dimethoxyanthracene-2-sulfonate, tris(4-t-butylphenyl)sulfonium trifluoromethanesulfonate, N-naphthalimide trifluoromethanesulfonate, p-nitrobenzyl 9,10-dimethoxyanthracene-2-sulfonate, 4-methoxy- α -({[(4-methylphenyl)sulfonyl]oxy}imino)benzeneacetonitrile, and 2-(2'-furylethenyl)-4,6-bis(trichloromethyl)-1,3,5-triazine. The salt is not necessarily limited to those listed above.

These sensitizers may be used individually or in combination. The amount of the sensitizer used is generally in the range of from 0.1 to 40% by weight, or the total amount of the two or more sensitizers used is preferably in the range of from 0.1 to 40% by weight, more preferably 1 to 20 parts by weight, relative to 100 parts by weight of polymer component (A).

As solvent (D) in the present invention, preferred is a polar solvent, such as γ -butyrolactone, N-methyl-2-pyrrolidone, N-acetyl-2-pyrrolidone, N,N-dimethylacetamide, N,N-dimethyl sulfoxide, hexamethylphosphoric triamide,

dimethylimidazolidinone, or N-acetyl- ϵ -caprolactam. Other than the polar solvent, ketone, ester, lactone, ether, halogenated hydrocarbon, or hydrocarbon, for example, acetone, diethyl ketone, methyl ethyl ketone, methyl isobutyl ketone, cyclohexanone, methyl acetate, ethyl acetate, butyl acetate, diethyl oxalate, diethyl malonate, diethyl ether, ethylene glycol dimethyl ether, diethylene glycol dimethyl ether, tetrahydrofuran, dichloromethane, 1,2-dichloroethane, 1,4-dichlorobutane, trichloroethane, chlorobenzene, o-dichlorobenzene, hexane, heptane, octane, benzene, toluene, or xylene may also be used. These organic solvents may be used individually or in combination. With respect to the type of the solvent, there is no particular limitation as long as the solvent dissolves the photosensitive resin composition of the present invention.

In addition to the essential components, the photosensitive resin composition of the present invention may also contain a silane coupling agent as an agent for improving the adhesion to a silicon substrate. R_2 in the formula (1) of the base polymer may be modified with diaminosiloxane. If desired, an addition-polymerizable compound, a dissolution inhibitor, a dissolution promoter, and a stabilizer may be incorporated.

A negative addition-polymerizable compound contains a reactive unsaturated functional group. Examples of the functional groups may include functional groups having an unsaturated double bond, such as a vinyl group, an allyl group, an acryloyl group, and a methacryloyl group, and functional groups having an unsaturated triple bond, such as propargyl group. Of these, preferred are functional groups of a conjugated type, such as a vinyl group, an acryloyl group, and a methacryloyl group, from the

viewpoint of achieving excellent reactivity. The number of the functional group or groups contained is preferably 1 to 4 from the viewpoint of obtaining excellent stability, and the functional groups contained may be either the same or different.

Examples of the addition-polymerizable compounds may include diethylene glycol diacrylate, triethylene glycol diacrylate, tetraethylene glycol diacrylate, diethylene glycol dimethacrylate, triethylene glycol dimethacrylate, tetraethylene glycol dimethacrylate, trimethylolpropane diacrylate, trimethylolpropane triacrylate, trimethylolpropane dimethacrylate, trimethylolpropane trimethacrylate, styrene, α -methylstyrene, 1,2-dihydronaphthalene, 1,3-diisopropenylbenzene, 3-methylstyrene, 4-methylstyrene, 2-vinylnaphthalene, butyl acrylate, butyl methacrylate, isobutyl acrylate, hexyl acrylate, isooctyl acrylate, isobornyl acrylate, cyclohexyl methacrylate, 1,3-butanediol diacrylate, 1,3-butanediol dimethacrylate, neopentyl glycol diacrylate, 1,4-butanediol diacrylate, 1,4-butanediol dimethacrylate, 1,6-hexanediol diacrylate, 1,6-hexanediol methacrylate, 1,9-nonanediol dimethacrylate, 1,10-decanediol dimethacrylate, dimethyloltricyclodecane diacrylate, pentaerythritol triacrylate, pentaerythritol tetraacrylate, pentaerythritol trimethacrylate, pentaerythritol tetramethacrylate, 2-hydroxyethyl acrylate, 2-hydroxyethyl methacrylate, 1,3-acryloyloxy-2-hydroxypropane, 1,3-methacryloyloxy-2-hydroxypropane, methylenebisacrylamide, N,N-dimethylacrylamide, and N-methylolacrylamide. These may be used individually or in combination.

The amount of the addition-polymerizable compound used is preferably 5 to 100% by weight, based on the weight of component (A), and is more preferably 5 to 40% by weight

from the viewpoint of achieving excellent compatibility. When the amount of the addition-polymerizable compound used is less than 5% by weight, the exposed portion tends to dissolve during the development and the film tends to be washed away after the development. When the amount is more than 100% by weight, the film tends to be washed away after the development, and the film being formed may be suffer whitening.

For improving the storage stability, a radical polymerization inhibitor or radical polymerization retarder may be added to the composition. Examples of the radical polymerization inhibitors or radical polymerization retarders may include p-methoxyphenol, diphenyl-p-benzoquinone, benzoquinone, hydroquinone, pyrogallol, phenothiazine, resorcinol, orthodinitrobenzene, paradinitrobenzene, metadinitrobenzene, phenanthraquinone, N-phenyl-1-naphthylamine, N-phenyl-2-naphthylamine, cupferron, phenothiazine, 2,5-toluquinone, tannic acid, parabenzylaminophenol, and nitrosoamine. These may be used individually or in combination. The amount of the radical polymerization inhibitor or radical polymerization retarder used is generally 0.01 to 30% by weight, based on the weight of component (A).

The photosensitive resin composition of the present invention may be applied to a substrate, such as a silicon wafer, a metallic substrate, or a ceramic substrate, by a dipping process, a spraying process, a screen printing process, or a spin coating process, and dried by appropriately heating for removing the solvent, to thereby form a tack-free film. This film is exposed to active rays of light or actinic radiation through a mask having a desired pattern. As the active rays of light or actinic radiation for exposure, a contact/proximity aligner, mirror

projection aligner, i-line stepper, or g-line stepper using an ultra high-pressure mercury lamp, ultraviolet light, a visible light source, an X-ray, or an electron beam may be used. Subsequently, if necessary, the resultant film is
5 subjected to post exposure bake (PEB) treatment, followed by development. When the photosensitive resin composition is negative, after the exposure, the unexposed portion may be removed by dissolving it in a developer to obtain a desired negative pattern. When the photosensitive resin
10 composition is positive, after the exposure, the exposed portion may be removed by dissolving it in a developer to obtain a desired positive pattern.

As the developer, an organic solvent or an alkaline aqueous solution may be used. Examples of the organic
15 solvent developer may include N-methyl-2-pyrrolidone, N,N-dimethylacetamide, N,N-dimethylformamide, dimethyl sulfoxide, tetramethylurea, hexamethylphosphoric triamide, γ -butyrolactone, water, alcohol, ketone, ester, lactone, ether, halogenated hydrocarbon, and hydrocarbon. Further
20 examples of the organic solvent may include methanol, ethanol, isopropyl alcohol, acetone, diethyl ketone, methyl ethyl ketone, methyl isobutyl ketone, cyclohexanone, methyl acetate, ethyl acetate, butyl acetate, diethyl oxalate, diethyl malonate, diethyl ether, ethylene glycol dimethyl
25 ether, diethylene glycol dimethyl ether, tetrahydrofuran, dichloromethane, 1,2-dichloroethane, 1,4-dichlorobutane, trichloroethane, chlorobenzene, o-dichlorobenzene, hexane, heptane, octane, benzene, toluene, and xylene. These may be used individually or in combination.

30 As the alkaline aqueous solution, there may be used an aqueous solution of an alkaline metal hydroxide, such as caustic potash or caustic soda, quaternary ammonium hydroxide, such as tetramethylammonium hydroxide,

tetraethylammonium hydroxide, or choline, or an aqueous solution of an amine, such as ethanolamine, propylamine, or ethylenediamine. After the development, if necessary, rinsing with water or a poor solvent is performed. As the
5 rinsing liquid, for example, methanol, ethanol, isopropyl alcohol, benzene, toluene, xylene, methyl cellosolve, or water is used. The thus obtained pattern is heated to remove the sensitizer and solvent to give a stable, high heat-resistant polyimide pattern.

10 The heating temperature is preferably 150 to 500°C, more preferably 200 to 400°C. When the heating temperature is lower than 150°C, the resultant film is likely to have poor mechanical properties and poor thermal properties, and, when the heating temperature is higher than 500°C, the
15 resultant film is likely to have poor mechanical properties and poor thermal properties.

The heating time is preferably 0.05 to 10 hours. When the heating time is shorter than 0.05 hour, the resultant film is likely to have poor mechanical properties and poor
20 thermal properties, and, when the heating time is longer than 10 hours, the resultant film is likely to have poor mechanical properties and poor thermal properties.

The photosensitive resin composition of the present invention may be used in an electronic part for
25 semiconductor device or multilayer wiring board. Specifically, the present photosensitive resin composition may be used in forming a surface protecting film layer or interlayer dielectric layer for semiconductor device, or an interlayer dielectric layer for multilayer wiring board.
30 The semiconductor device directed to the present invention is not particularly limited as long as it has a surface protecting film layer or interlayer dielectric layer formed

of the composition of the present invention, and it may be of various structures.

An example of the process for fabricating a semiconductor device as one example of the electronic part of the present invention is described below.

Fig. 1 is a manufacturing process diagram of a semiconductor device of a multilayer wiring structure. In Fig. 1, a semiconductor substrate 1, such as an Si substrate, having a circuit element is covered with a protecting film 2, such as a silicon oxide film, excluding the circuit element at a predetermined portion. On the exposed circuit element, there is formed a first conductive layer 3. An interlayer dielectric layer 4 is formed on the semiconductor substrate by, e.g., a spin coating process (step (a)).

A photosensitive resin layer 5 made of, e.g., chlorinated rubber or phenolic novolak is then formed on the interlayer dielectric layer 4 by a spin coating process, and windows 6A are formed by a known photolithography technique so that predetermined portions of the interlayer dielectric layer 4 are exposed (step (b)).

The interlayer dielectric layer 4 exposed through the windows 6A is selectively etched by a dry etching method using gas, such as oxygen gas or carbon tetrafluoride gas, to form windows 6B. The photosensitive resin layer 5 is then completely removed using such an etching solution that only the photosensitive resin layer 5 is etched without etching the first conductive layer 3 exposed through the windows 6B (step (c)).

A second conductive layer 7 is formed further using a known photolithography technique to complete electrical connection to the first conductive layer 3 (step (d)).

When a multilayer wiring structure having three layers or more is formed, a series of the aforementioned steps is repeated to form layers.

Next, a surface protecting film layer 8 is formed. In the example of Fig. 1, the surface protecting film layer 8 is formed by applying the photosensitive resin composition by a spin coating process and drying it, and exposing the resultant layer to light through a mask having a pattern for forming windows 6C at predetermined portions, and then developing it using an alkaline aqueous solution to form a pattern, and heating the layer to obtain a resin film. This resin film protects the conductive layer from an external stress or α -line, and therefore the semiconductor device obtained has excellent reliability. In the aforementioned example, the interlayer dielectric layer 4 may also be formed of the photosensitive resin composition of the present invention.

Examples

The present invention will be explained hereinbelow with reference to the examples.

Synthesis Example 1

While stirring, 2.34 g (0.018 mol) of 2-hydroxyethyl methacrylate was added to a solution consisting of 100 ml of dried N-methylpyrrolidone and 21.18 g (0.072 mol) of 3,3',4,4'-biphenyltetracarboxylic dianhydride in a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, and the resultant mixture was stirred at 70°C for one hour, subsequently at room temperature overnight. To the resultant reaction solution was added dropwise over one hour a solution of 6.49 g (0.06 mol) of metaphenylenediamine and 0.25 g (0.001 mol) of 1,3-

bis(3-aminopropyl)tetramethyldisiloxane in 80 ml of dried N-methylpyrrolidone in a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, followed by stirring at room temperature overnight. While stirring, to the resultant reaction solution was then added dropwise over one hour a solution of 29.92 g (0.145 mol) of N,N-dicyclohexylcarbodiimide in 100 ml of dried N-methylpyrrolidone. To the reaction solution was added 45.5 g (0.35 mol) of 2-hydroxyethyl methacrylate, and the resultant mixture was stirred at 50°C for 5 hours, and subsequently at room temperature overnight. The resultant reaction mixture was diluted with 50 ml of acetone and unnecessary substances were removed from the mixture by suction filtration, and the resultant filtrate was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, subsequently with methanol, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain polymer P-1.

Synthesis Example 2

In a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution consisting of 80 ml of dried N-methylpyrrolidone, 14.4 g (0.072 mol) of 4,4'-oxydianiline, and 0.63 g (0.008 mol) of pyridine was cooled to 0°C. To the solution was added dropwise 1.12 g (0.008 mol) of benzoyl chloride, followed by stirring at room temperature for 30 minutes, to obtain reaction solution A-1. 100 ml of dried N-methylpyrrolidone and 21.18 g (0.072 mol) of 3,3',4,4'-biphenyltetracarboxylic dianhydride were then added to another closed reaction vessel having a stirrer, a

thermometer, and a nitrogen gas feed pipe, and, while stirring, the above-obtained reaction solution A-1 was added dropwise over 30 minutes. The resultant mixture was stirred at 50°C for 5 hours, subsequently at room
5 temperature overnight. The resultant reaction solution was cooled to 0°C, and 10.4 g (0.11 mol) of chloromethyl ethyl ether and 11.1 g (0.11 mol) of triethylamine were successively added dropwise to the solution over 30 minutes, followed by stirring at room temperature for 30
10 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content
15 thereof became less than 1.0% by weight at room temperature to obtain polymer P-2.

Synthesis Example 3

31.0 g (0.1 mol) of bis(3,4-dicarboxyphenyl) ether
20 dianhydride was added to 100 ml of dried N-methylpyrrolidone and 14.8 g (0.2 mol) of n-butanol in a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, and the resultant mixture was stirred at 70°C for 24 hours to obtain an ester. The
25 resultant solution was cooled to 0°C, and 23.8 g (0.2 mol) of thionyl chloride was added dropwise to the solution, followed by stirring for 30 minutes, to obtain reaction solution A-2. 100 ml of dried N-methylpyrrolidone, 33.0 g (0.09 mol) of 2,2-bis(3-amino-4-hydroxyphenyl)-1,1,1,3,3,3-
30 hexafluoropropane, and 15.82 g (0.2 mol) of pyridine were then added to another closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, and

the solids therein were dissolved by stirring. The resultant solution was cooled to 0°C, and the above-obtained reaction solution A-2 was added dropwise to the solution over 30 minutes, followed by stirring at room temperature for 30 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain polymer P-3.

Synthesis Example 4

In a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution consisting of 100 ml of dried N-methylpyrrolidone and 25.8 g (0.1 mol) of 4,4'-oxybis(carboxyphenyl) was cooled to 0°C, and 23.8 g (0.2 mol) of thionyl chloride was added dropwise to the solution, followed by stirring for 30 minutes, to obtain reaction solution A-3. In another closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution consisting of 100 ml of dried N-methylpyrrolidone, 36.6 g (0.1 mol) of 2,2-bis(3-amino-4-hydroxyphenyl)-1,1,1,3,3,3-hexafluoropropane, and 0.79 g (0.01 mol) of pyridine was cooled to 0°C, and 1.57 g (0.01 mol) of phenyl chloroformate was added dropwise to the solution, followed by stirring at room temperature for 30 minutes. 15.82 g (0.2 mol) of pyridine was added to the resultant mixture and cooled to 0°C, and then the above-obtained reaction solution A-3 was added dropwise to the mixture over 30 minutes, followed by stirring at room temperature for 30 minutes. The resultant

reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum
5 drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain polymer P-4.

Synthesis Example 5

In a closed reaction vessel having a stirrer, a
10 thermometer, and a nitrogen gas feed pipe, a solution consisting of 100 ml of dried N-methylpyrrolidone and 9.76 g (0.05 mol) of 1-methyl 2-aminoterephthalate was cooled to 0°C, and 8.85 g (0.05 mol) of benzenesulfonyl chloride was added dropwise to the solution, followed by stirring for 30
15 minutes. 15.82 g (0.2 mol) of pyridine was then added to the resultant mixture and stirred at room temperature for 30 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged
20 water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain chain extender B-1 (for the sake of simplicity, component (B) is referred to hereinafter as "chain
25 extender").

Synthesis Example 6

In a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution
30 consisting of 100 ml of dried N-methylpyrrolidone and 18.3 g (0.05 mol) of 2,2-bis(3-amino-4-hydroxyphenyl)-1,1,1,3,3,3-hexafluoropropane was cooled to 0°C, and 12.1 g (0.1 mol) of pivaloyl chloride was added dropwise to the

solution, followed by stirring for 30 minutes. 15.82 g (0.2 mol) of pyridine was then added to the resultant mixture and stirred at room temperature for 30 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain chain extender B-2.

Synthesis Example 7

In a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution consisting of 100 ml of dried N-methylpyrrolidone and 7.6 g (0.05 mol) of 3,5-diaminobenzoic acid was cooled to 0°C, and 15.7 g (0.1 mol) of phenyl chloroformate was added dropwise to the solution, followed by stirring for 30 minutes. 15.82 g (0.2 mol) of pyridine was then added to the resultant mixture and stirred at room temperature for 30 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain chain extender B-3.

Synthesis Example 8

In a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution consisting of 100 ml of dried N-methylpyrrolidone and 25.9 g (0.05 mol) of 4'',4''',-(hexafluoroisopropylidene)-bis(4-

phenoxyaniline) was cooled to 0°C, and 16.4 g (0.1 mol) of diisopropylcarbamyl chloride was added dropwise to the solution, followed by stirring for 30 minutes. 15.82 g (0.2 mol) of pyridine was then added to the resultant mixture and stirred at room temperature for 30 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain chain extender B-4.

Synthesis Example 9

In a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe, a solution consisting of 100 ml of dried N-methylpyrrolidone and 6.86 g (0.05 mol) of 4-aminobenzoic acid was cooled to 0°C, and 3.93 g (0.05 mol) of acetyl chloride was added dropwise to the solution, followed by stirring for 30 minutes. 15.82 g (0.2 mol) of pyridine was then added to the resultant mixture and stirred at room temperature for 30 minutes. The resultant reaction mixture was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain chain extender B-5.

Synthesis Example 10

While cooling, 0.3 mol of triethylamine was added dropwise to a solution of 0.1 mol of tris(4-

hydroxyphenyl)methane and 0.29 mol of naphthoquinone-1,2-diazido-4-sulfonyl chloride in 150 g of dioxane in a closed reaction vessel having a stirrer, a thermometer, and a nitrogen gas feed pipe to effect a reaction, and the resultant reaction mixture was subjected to filtration, and the filtrate was treated by vigorously stirring in 2.0 L of ion-exchanged water. The solid precipitate was further washed with ion-exchanged water, and subjected to suction drying on a filter and then subjected to vacuum drying until the moisture content thereof became less than 1.0% by weight at room temperature to obtain orthoquinonediazide compound C-1.

Example 1

100 g of P-1 obtained in the Synthesis Example 1, 140 g of N-methylpyrrolidone, and 0.3 g of p-methoxyphenol were placed in a three-necked flask having a stirrer, a thermometer, and a nitrogen gas feed pipe, and the solids therein were dissolved by stirring. To the resultant solution were added 6.0 g of 2,2'-bis(o-chlorophenyl)-4,4',5,5'-tetraphenylbiimidazole, 3.0 g of 2-mercaptobenzoxazole, 0.6 g of ethyl Michler's ketone as a sensitizer, 18.0 g of tetraethylene glycol diacrylate as an addition-polymerizable compound, and 5.0 g of chain extender B-1 obtained in the Synthesis Example, and the solids therein were dissolved by stirring at room temperature for about 24 hours, followed by filtration using a filter, to obtain a photosensitive resin composition solution. The solution obtained was applied onto a 5-inch silicon wafer by spin coating and then dried to form a 5.0 ± 1.0 μm thick film. The film was then subjected to exposure with a pattern mask using an I-line stepper at a light exposure of 200 to 1000 mJ/cm². The

resultant wafer was allowed to stand in a light shielded box for one hour, and then subjected to paddle development using γ -butyrolactone, thus obtaining an excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . The
5 aforementioned solution was applied onto a 5-inch silicon wafer by spin coating and then dried to form an $18.0 \pm 1.0 \mu\text{m}$ thick film, and then, the film was subjected to exposure with a pattern mask using a contact aligner at a light
10 exposure of 1000 mJ/cm^2 . The resultant wafer was allowed to stand in a light shielded box for one hour, and then subjected to paddle development using γ -butyrolactone to obtain a 10 mm \times 120 mm strip pattern. The wafer was baked for cure in an oven filled with nitrogen gas at 320°C for
15 one hour to obtain a cured film. The strip thin film was peeled off the silicon wafer using an aqueous solution of hydrofluoric acid and dried, and then an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 50%.

20

Example 2

The same procedure as in the Example 1 was repeated except that 5.0 g of chain extender B-2 obtained in the Synthesis Example 6 was used to give an excellent relief
25 pattern having a resolution as high as 15 μm (minimum line width) at a light exposure of 450 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 1, and an elongation of the strip thin film was measured using an autograph. It was found that the
30 elongation was 70%.

Example 3

The same procedure as in the Example 1 was repeated except that 5.0 g of chain extender B-3 obtained in the Synthesis Example 7 was used to give an excellent relief pattern having a resolution as high as 15 μm (minimum line width) at a light exposure of 450 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 1, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 55%.

10

Example 4

The same procedure as in the Example 1 was repeated except that 5.0 g of chain extender B-4 obtained in the Synthesis Example 8 was used to give an excellent relief pattern having a resolution as high as 15 μm (minimum line width) at a light exposure of 500 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 1, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 55%.

20

Example 5

100 g of P-2 obtained in the Synthesis Example 2 and 140 g of N-methylpyrrolidone were placed in a three-necked flask having a stirrer, a thermometer, and a nitrogen gas feed pipe, and the solids therein were dissolved by stirring. To the resultant solution were added 5.0 g of p-nitrobenzyl-9,10-dimethoxyanthracene-2-sulfonate and 5.0 g of chain extender B-1 obtained in the Synthesis Example 5, and the solids therein were dissolved by stirring at room temperature for about 24 hours, followed by filtration using a filter, to obtain a photosensitive resin composition solution. The solution was treated in

30

accordance with the same procedure as in the Example 1 except that, after the exposure, the wafer was heated at 120°C for 60 seconds by a hot plate and then subjected to paddle development using a 2.38% aqueous solution of
5 tetramethylammonium hydroxide, to give an excellent relief pattern having a resolution as high as 15 μm (minimum line width) at a light exposure of 800 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 1 except that, after the exposure, the wafer
10 was heated at 120°C for 60 seconds by a hot plate and then subjected to paddle development using a 2.38% aqueous solution of tetramethylammonium hydroxide, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 40%.

15

Example 6

The same procedure as in the Example 5 was repeated except that 5.0 g of chain extender B-2 obtained in the Synthesis Example 6 was used to give an excellent relief
20 pattern having a resolution as high as 20 μm (minimum line width) at a light exposure of 900 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 5, and an elongation of the strip thin film was measured using an autograph. It was found that the
25 elongation was 40%.

Example 7

The same procedure as in the Example 5 was repeated except that 5.0 g of chain extender B-3 obtained in the Synthesis Example 7 was used to give an excellent relief
30 pattern having a resolution as high as 20 μm (minimum line width) at a light exposure of 900 mJ/cm^2 . A strip thin

film was obtained in accordance with the same procedure as in the Example 5, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 45%.

5

Example 8

100 g of P-3 obtained in the Synthesis Example 3 and 140 g of N-methylpyrrolidone were placed in a three-necked flask having a stirrer, a thermometer, and a nitrogen gas feed pipe, and the solids therein were dissolved by stirring. To the resultant solution were added 6.0 g of sensitizer C-1 obtained in the Synthesis Example 10 and 5.0 g of chain extender B-1 obtained in the Synthesis Example 5, and the solids therein were dissolved by stirring at room temperature for about 24 hours, followed by filtration using a filter, to obtain a photosensitive resin composition solution. The solution was treated in accordance with the same procedure as in the Example 1 except that a 2.38% aqueous solution of tetramethylammonium hydroxide was used in the paddle development to give an excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm². A strip thin film was obtained in accordance with the same procedure as in the Example 1 except that a 2.38% aqueous solution of tetramethylammonium hydroxide was used in the paddle development, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 40%.

30 Example 9

The same procedure as in the Example 8 was repeated except that 5.0 g of chain extender B-3 obtained in the Synthesis Example 7 was used as a chain extender to give an

excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 8, and an elongation of the
5 strip thin film was measured using an autograph. It was found that the elongation was 50%.

Example 10

The same procedure as in the Example 8 was repeated
10 except that 5.0 g of chain extender B-4 obtained in the Synthesis Example 8 was used as a chain extender to give an excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . A strip thin film was obtained in accordance with the same
15 procedure as in the Example 8, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 50%.

Example 11

20 100 g of P-4 obtained in the Synthesis Example 4 and 140 g of N-methylpyrrolidone were placed in a three-necked flask having a stirrer, a thermometer, and a nitrogen gas feed pipe, and the solids therein were dissolved by stirring. To the resultant solution were added 6.0 g of
25 sensitizer C-1 obtained in the Synthesis Example 10 and 5.0 g of chain extender B-4 obtained in the Synthesis Example 8, and the solids therein were dissolved by stirring at room temperature for about 24 hours, followed by filtration using a filter, to obtain a photosensitive resin
30 composition solution. The solution was treated in accordance with the same procedure as in the Example 1 except that a 2.38% aqueous solution of tetramethylammonium hydroxide was used in the paddle development to give an

excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 350 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 1 except that a 2.38% aqueous solution of tetramethylammonium hydroxide was used in the paddle development, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 70%.

10 Example 12

The same procedure as in the Example 11 was repeated except that 5.0 g of chain extender B-5 obtained in the Synthesis Example 9 was used to give an excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 11, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 55%.

20

Comparative Example 1

The same procedure as in the Example 1 was repeated except that chain extender B-1 obtained in the Synthesis Example 5 was not used to give an excellent relief pattern having a resolution as high as 15 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 1, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 30%.

30

Comparative Example 2

The same procedure as in the Example 4 was repeated except that chain extender B-2 obtained in the Synthesis Example 6 was not used to give an excellent relief pattern having a resolution as high as 20 μm (minimum line width) at a light exposure of 900 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 4, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 15%.

10

Comparative Example 3

The same procedure as in the Example 6 was repeated except that chain extender B-1 obtained in the Synthesis Example 5 was not used to give an excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 6, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 10%.

20

Comparative Example 4

The same procedure as in the Example 8 was repeated except that chain extender B-4 obtained in the Synthesis Example 8 was not used to give an excellent relief pattern having a resolution as high as 10 μm (minimum line width) at a light exposure of 400 mJ/cm^2 . A strip thin film was obtained in accordance with the same procedure as in the Example 8, and an elongation of the strip thin film was measured using an autograph. It was found that the elongation was 30%.

30

The combinations of the photosensitive polymers and chain extenders, the photosensitive properties, and the results of the measurement of elongation are shown in Table 1 below.

5

Table 1

No.	Polymer	Chain extender	Sensitivity (mJ/cm ²)	Resolution (μm)	Elongation (%)
Example 1	P-1	B-1	400	10	50
Example 2	P-1	B-2	450	15	70
Example 3	P-1	B-3	450	15	55
Example 4	P-1	B-4	500	15	55
Example 5	P-2	B-1	800	15	40
Example 6	P-2	B-2	900	20	40
Example 7	P-2	B-3	900	20	45
Example 8	P-3	B-1	400	10	40
Example 9	P-3	B-3	400	10	50
Example 10	P-3	B-4	400	10	50
Example 11	P-4	B-4	350	10	70
Example 12	P-4	B-5	400	10	55
Comparative Example 1	P-1	None	400	15	30
Comparative Example 2	P-2	None	900	20	15
Comparative Example 3	P-3	None	400	10	10
Comparative Example 4	P-4	None	400	10	30

INDUSTRIAL APPLICABILITY

10 The present invention provides a resin composition comprising a photosensitive polyimide precursor, polybenzoxazole precursor, polyimide, or polybenzoxazole capable of forming a cured resin having excellent properties, a copolymer thereof, or a mixture thereof; a method for forming a pattern using the resin composition; and an electronic part having an electronic device having the pattern.